

PHARMACEUTICAL COMPOSITIONS COMPRISING
NATURAL HUMAN α -INTERFERON

- This application is a United States national stage application based on
- 5 International Application No. PCT/IT97/00040, filed February 27, 1997, which claims priority to Italian Patent Applications Nos. RM96A000136 filed February 28, 1996 and RM96A000247 filed June 14, 1996.

[0001] The invention concerns pharmaceutical compositions for a peroral

10 administration comprising natural human α -interferon isolated from lymphoblastoid or leukocytic [leukocytic] cells. In particular compositions are useful for therapy of viral infections, in particular viral hepatitis, neoplasia and immunodeficiency syndromes. The effective doses of interferon [efficient dosages] are clearly lower than dosages utilized for parenteral administration.

15 [0002] α -, β -, γ - interferons are usually administered by injection and are used for therapy. α - interferon is the most largely utilized interferon (1). In an updated study of medicaments for either acute or chronic viral hepatitis therapy (2), only α -interferon is widely accepted as a single therapeutic agent.

[0003] "Viral hepatitis" means at least five different pathologies, having different

20 causative agents [, namely] and designated hepatitis A, B, C, D, or E.

[0004] The therapeutic trend is to treat said pathologies with α -interferon, with dosages according to the kind of hepatitis, to the overall status of the subject and to other variable factors. In general, further to the interferon treatment an almost [normalisation] normalization in clinical and biochemical parameters is achieved for chronic hepatitis

25 (B,C, or D). The effect of interferon [activity] on acute hepatitis has not been [focused] determined yet, although [though] for hepatitis C, a therapeutic treatment with α -interferon lowers the [chronicitation rate] chronicity of the disease.

[0005] Therapeutic cycles [indicate the day] consist of alternate day subcutaneous administration [through subcutaneous route] of recombinant α -interferon (r

30 α -IFN) at dosages of [app. 5.000.000] approximately 5,000,000 UI, that in special cases can be up to [9.000.000] 2,000,000 UI/day.

[0006] The length of therapeutic cycles is [of] from six months up to one year (nine months average).

[0007] In many cases, undesired side effects interfere with the course of therapeutic treatment. In fact some patients, in particular those at an advanced stage of disease or with severe physiologic [damages] damage, do not tolerate the therapy and therefore the treatment should be interrupted. Claimed side effects are: fever, nausea, 5 [vomit] vomiting, tiredness, [algia] pain and depression.

[0008] Moreover the therapeutic [cost] costs are quite relevant due to the high amount of active principle (more than [8.000] 8,000 new cases each year in Italy and [300.000] 300,000 world-wide) and to the necessity of [hospitalisation] hospitalization just in consideration of said side effects further to the parenteral administration (day 10 hospital or outpatient's department).

[0009] Finally, as far as chronic active viral hepatitis the only alternative to the interferon treatment is [represented by] liver transplant.

[0010] The clinical trend is to increase the [posology] dosage and the length of therapeutic cycle (3), but clinical data show (4): sever side effects; low acceptance by the 15 patient; and high therapeutic costs. Garcia et al. (5) report that the estimate for each cured patient is between [700.000 and 2.000.000] 700,000 and 2,000,000 English pounds. Capri S. (6) report that the cost of each interferon treatment is [of Lit. 70.000.000] 70,000,000 Lire/subject.

[0011] It is therefore evident that the actual composition of interferon for 20 therapeutic treatment of hepatitis is not optimal.

[0012] Moreover clinical results show a better therapeutic efficacy in patients which are not the main target for therapy, namely: young subjects, subjects with a disease at an initial stage, subjects infected with [genotypic] genotypic virus types 2 or 3, and subjects having low viremia [subjects]. [On the contrary a] In contrast, less therapeutic 25 efficacy can be found in those subjects which really need the therapeutic treatment (subjects [poco] less respondent), as subjects affected by an aggressive form (active chronic hepatitis) [,long length diseases affected subjects, over 50 subjects] or by long-term disease. Thus patients that really need an immediate interferon treatment are those that have a lower chance of [success] being treated successfully.

30 [0013] The authors of the instant invention have found a pharmaceutical composition comprising natural human α -interferon from either lymphoblastoid or

[leukocitic] leukocytic cells to be administered [through] by the peroral route, with dosages clearly lower than those used for parenteral administration. The composition maintains [as] unaltered the chemical-physical, biological and pharmacological characteristics of the active principle, having a therapeutic effect substantially analogous to the compositions of the prior art but overcoming its disadvantages [thereof].

[0014] The composition is preferably in a liquid form with a concentration of 100 to 500 IU/ml, preferably [approx.] approximately 150 UI/ml, most preferably in [mono-dosage] single-dosage units, most preferably of [appr.] approximately 1 ml.

[0015] The composition acts by activating the [defence] defense mechanisms against viral infections, [tumour] tumor growth and [stimulates] by stimulating an immune response.

[0016] The [utilisation] utilization of natural interferon was chosen for the better chances of therapeutic success with respect [respects] to recombinant interferon, obtained by cloning of a single subtype.

[0017] Though [leukocitic] leukocytic and lymphoblastoid interferons exert the same therapeutic properties, the former can be advantageously produced. As a matter of fact it is obtainable by [stabilised] stabilized cell lines, without the need of blood donors.

[0018] Processes for purifying interferons are known to those skilled in the art, and for example are shown in US Patent 4,732,683; in Cantell K. and Hirvonen S. Texas Reports on Biology and Medicine, Vol. 35, p. 138, 1977; and in Zoon K.C. et al., Science 207, p. 527, 1980.

[0019] The peroral route is generally much more accepted by subjects, makes easier [posology] therapeutic schemes and dosages, lowers to stops the antigenic risk, induces the transmission and amplification signal mechanism, with a [mirato] therapeutic effect, with dosages 100 times lower than known formulations for parenteral administrations.

[0020] The low dosage annuls the risk of toxic effects; allows a better availability of medicine to satisfy an increasing request and a drastic lowering of therapeutic costs.

[0021] The preferred formulation in dosage units of small volumes (1 ml) to drink allows an immediate availability of the active principle, a good standard of cleanliness from the [monodosage] single dose primary container; the certainty of the

taken dosage; the taking of the active principle to be immediately absorbed by the oropharyngeal mucosa, easily preventing the deglutition, an [ease] easy and safe way of administration for all patients, [as opposite to] as compared to lozenges or [tablets] tablet formulations that should be kept in the mouth [till to full dissolution] until completely dissolved, with high chances [of swallowing] that the formulation will be swallowed.

[0022] Moreover the composition of the invention is conveniently used for home therapies or [on] at the work place, as a precautionary measure for the prophylaxis of viral pathologies, and to control chronic diseases which [need of long therapeutic cycles (even yearly) and often recurrent] require long (even yearly) and often recurrent therapeutic cycles.

[0023] The composition can be used also in association with other drugs to get synergism and optimize therapeutic schemes.

[0024] The following clinical studies show the therapeutic effect. A comparison of the electrophoretic protein pattern and of the concentration of IgG, IgA, and IgM, before the beginning of the peroral therapy with natural human α -interferon of hepatitis or other pathologies [affected subjects,] before and after two weeks of therapeutic treatment, allows [to foreseen quali-quantitatively] predicting semi-quantitatively the subject response.

[0025] Subjects which respond to the therapy with 450 UI/[die]day dosages show a decrease of α 2- and β -globulins, of IgGs, of the IgG/IgA ration, together [to] with an increase of IgA and IgM concentrations, have a good chance of [eliminate] eliminating the HBVe antigen and to seroconvert, namely to confer a stable remission of the pathology.

[0026] On the other hand subjects which respond to the same therapy with a decrease of [albumin] serum albumin concentration, of IgGs, IgAs, IgMs, together [to] with an increase of α 1-globulin fractions, should seroconvert with longer times.

[0027] Moreover subjects that respond with an increase of IgGs, of the IgG/IgA ration, together [to] with a decrease of IgM and of the IgA/IgM [ration] ratio, could be resistant to the therapy.

[0028] The monitoring of said parameters (markers) is useful for a planning of therapeutic strategies in clinic and also for the clinical practitioner.

[0029] Clinical studies on healthy subjects.

Table 1 shows different therapeutic schemes.

Table 1

5	Exp.		active comp.	No. admin./day	Dosages	days trt.	blood bleedings
10	A	aA	α -IF	1 (3dsg)	450 UI	1	T ₀ , T ₁ , T ₂ , T ₃
		aB	placebo	1 (3dsg)	-	1	T ₀ , T ₁ , T ₂ , T ₃
	B	bA	α -IF	1 (3dsg)	450 UI	5	T ₀ , T ₁ , T ₂ , T ₃ , T ₄ , T ₅ , T ₆ , T ₇
		bB	placebo	1 (3dsg)	-	5	T ₀ , T ₁ , T ₂ , T ₃ , T ₄ , T ₅ , T ₆ , T ₇
15	C	cA ₁	α -IF	2 (1 dsg)	300 UI	1	T ₀ , T ₁ , T ₂ , T ₃
		cA ₂	α -IF	3 (1dsg)	450 UI	1	T ₀ , T ₁ , T ₂ , T ₃
		cb	placebo	3 (1dsg)	-	1	T ₀ , T ₁ , T ₂ , T ₃
	D	dA ₁	α -IF	2 (1dsg)	300 UI	5	T ₀ , T ₁ , T ₂ , T ₃ , T ₄ , T ₅ , T ₆ , T ₇
20		dA ₂	α -IF	3 (1dsg)	450 UI	5	T ₀ , T ₁ , T ₂ , T ₃ , T ₄ , T ₅ , T ₆ , T ₇
		dB	placebo	3 (1dsg)	-	5	T ₀ , T ₁ , T ₂ , T ₃ , T ₄ , T ₅ , T ₆ , T ₇

wherein T₀ = background; T₁ = 1 d [further] after the first administration, T₂ = 2 d

25 [further] after the first administration, T₃ = 3d [further] after the first administration, T₄ =
4d [further] after the first administration, T₅ = 5d [further] after the first administration,
T₆ = 1d after the treatment suspension, and T₇ = 2d after the treatment suspension.

[0030] The change of the induced biological response with respect to the
therapeutic scheme, has been measured on samples of blood, taken at different times. In
30 particular the activity with respect to the day dosage of active principle, to the mono- or
pluri-administration, and to the length of the therapeutic cycle was measured.

[0031] The analysis of data show that natural human α -interferon from either lymphoblastoid or [leukocitic] leukocytic cells, administered at low dosages for a peroral route, is able to modulate (according to the dosage and to the length of the therapeutic cycle) the expression of membrane antigen of healthy subject blood mononuclear cells.

5 In particular, according to the therapeutic scheme, the pharmaceutical composition seems to be able to increase both CD4 and CD8 cell [population] populations. [It is also evident an increased] Increased expression of markers of cell activation, such as DR antigens and the interleukin 2 receptor, is also evident.

[0032] The therapeutic scheme with 450 U/[die]day x 5 d (exp. b) [is the one] 10 provided better results, as shown in Tables 2 and 3. In fact there is an increase (% and absolute) [of] in CD3, CD4, DR1, and CD25 lymphocytes. said increases are, according to different cases, [better] more evident at T₃, T₄, and T₅ times [to] and later decrease at T₆ and T₇ times.

[0033] The same [posology] dosage, but with a shorter therapeutic cycle (1 day) 15 (exp.a), interferes less evidently with the [%] percentage and absolute numbers of mononuclear cells in the blood (Tables 4 [e] and 5). In fact in this experiment an increase of average percentage values but not of absolute T, CD8, and class I [hystocompatibility] histocompatibility antigen [lymphocytes] lymphocyte values, is evident at time T₃.

[0034] Other experimental conditions show lower increases of the immune 20 response.

[0035] Therefore, natural human α -interferon from either lymphoblastoid or [leukocitic] leukocytic cells, administered at low dosages [trough] via the peroral route, [shows] plays an important role in modulating the immune response, both in the afferent and efferent phases [phase afferent than efferent, e] and has a therapeutic application for 25 the treatment of infective diseases and of other conditions of immunodeficiency.

Clinical studies on hepatitis subjects.

Viral B Hepatitis

[0036] 14 patients affected by chronic viral B hepatitis, [with an aged comprised] between 4 and 59 years of age, were used for random studies.

30 [0037] All [of subject] subjects were previously treated for different periods ranging from some months to some years with steroids or with steroid-azothiopurine,

with no beneficial effects, neither for the clinical symptomatology nor for the biochemical parameters of the disease, which evolved, in some cases, to hepatic cirrhosis.

[0038] The therapeutic treatment of [a] one administration of 150 U/day was initiated immediately after the suspension of the previous treatment, and effects of said
5 treatment were monitored by checking any alteration of the immune response; of the haematological and biochemical parameters; of serum markers of the viral infection and of the [hystochemistry] histochemistry of hepatic [bioptic] biopsy samples.

[0039] The time of observation varied from 15 to 32 months and the results can be summarized [in the following] as follows:

10 1) all [of] patients during the first 3-6 weeks of treatment registered a transient decay of hepatic biochemical functions (i.e. a 2-3 fold increase of alanine aminotransferase (ALT) levels), with no clinical symptoms of disease worsening;

2) the phenomenon [goes on] continued for 4-6 weeks;

15 3) in all [of] treated patients an intense activation of the immune response was observed, even after the therapeutic treatment;

4) 7 patients [eliminate] eliminated HBV DNA and HBeAg from their serum and were [stable seroconvert] stably seroconverted;

5) 1 patient [has] had an [HBcAg increased title] increased HBeAg titer, more than the original value; and

20 6) in the other 9 patients said [titre decreases] titer decreased significantly.

[0040] Therefore, 50% of patients get a stable remission of the disease.

Viral C Hepatitis

[0041] The therapeutic standard of viral hepatitis C foresees the use of α -interferon through the parenteral route.

25 [0042] 6 active chronic hepatitis C affected patients were subjected to therapy with peroral administration at 150U/[die]day, by starting the treatment just after the suspension of the steroid therapy.

[0043] The observation time (equal to the length of the treatment) [resulted to be variable] varied from 19 to 69 weeks. In general the treatment was well tolerated and all
30 of the patients registered a significant increase of vivacity and appetite, with a better tolerance to physical exercises.

[0044] No patients got a normalization of transaminase levels during the observation period, but one [which registered] demonstrated the biochemical and clinical remission of the disease, after the treatment suspension at the 19th week due to an [increasing] increase in articular pains.

5 [0045] Results are shown in tables 2-5.

[0046] BIBLIOGRAPHY

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